

JERZEJEWSKA, AICJA

Poland/Pharmacology. Toxicology. Chemo-Therapeutical Pre- U-7
parations.

Abs Jour : Ref Zhur-Biol., No 7, 1958, 33052

Author : Kossowski Stanislaw, Bekierkunst Adam, Agopsowicz
Grzegorz, Jerzejewska Aicja.

Inst : Not given

Title : Therapy of Azaena with Dihydrostreptomycin and
a Mixture of Dihydrostreptomycin and Penicillin.

Orig Pub : Arch. immunol. i terap. doswiadcz., 1955, 3,
239-247

Abstract : Twenty-three patients ill with azaena were trea-
ted with dihydrostreptomycin (I); 30 other patients
were given dihydrostreptomycin and penicillin (II)
simultaneously. The patients of the 1st group were
administered 1 in doses of 0.5 to 1g every 24
hours for a period of 12 days. Those of the 2nd group

Card 1/2

FIALOVA, V.; JERZEK, V., MIKULENKA, V.

The relationship of pulse velocity to blood lipids in atherosclerosis.
Cor Vasa 4 no.1:20-25 '62.

1. The IIInd Internal Clinic, Charles University, Prague.
(LIPIDS blood) (ARTERIOSCLEROSIS physiol)
(PULSE physiol)

JERZMANOWSKA, Zofia, prof. dr

Certain development trends of organic chemistry. Problemy
20 no. 4: 197-204 '64.

1. Kierowniczka Katedry Chemii Organicznej, Wydział Farmace-
utyczny, Akademia Medyczna, Łódź.

Hyperin, a glucoside of *Hypericum perforatum* L. *Zola, Jagiennowska, Wladimirov farm.* 64, 627-32 (1937); *Chem. Zentr.* 1938, 1, 331.—A new glucoside, hyperin (I), was isolated from the com. drug herba *hyperici concisa sicca*. The drug (330 g.) was extd. 3 hrs. with 1 l. of 80% alc., then 3 hrs. with 800 cc. alc. The ext. was evapd. to 150 cc. in *vacuo* and, after the addn. of water, freed from chlorophyll with CS₂. The alc. soln. was treated with 2 vols. of ether, which caused a tarry liquid to sep. The ether-alc. soln. was then evapd. in *vacuo* to 50 cc. at a temp. not exceeding 40° and the evap. completed in a vacuum desiccator. When

the residue of 8.12 g. was ground with acetone & a grayish yellow ppt. of the raw prep. The yield of the crude glucoside was 0.6-0.7% (m. p. 210-13°). The prep., when recrystd. from water, was light yellow. It is sol. in 200 parts water and gives pale yellow needles with 1.5, 2, 3 and 3.5 mols. water. The last hydrate is most frequently obtained. The hydrates are stable in the air but the anhyd. prep. is very highly hygroscopic. The pure compd. m. 217-8° (decomps. 1, [α]_D 25° soln in pyridine-alk. is -50° & 0°). It is sol. in the cold only in pyridine, has an acid character, gives a brownish green color with FeCl₃, a yellow ppt. with Pb acetate, reduces an NH₃-AgNO₃ soln. and reduces Fehling soln. slowly when warmed. It gives a violet Molisch reaction. The anhyd. glucoside has the formula C₁₂H₁₅O₆. Hydrolysis of the glucoside with dil. acids yields quercetin. The hexose combined with the quercetin was shown to be D-galactose. I is therefore quercetin-β-galactoside. By methylation with CH₃N₃ it was shown that I is the 3-O-galactoside. A 2-g. sample of I was allowed to stand 24 hrs. in a mixt. of 100 cc. MeOH and 1.00 cc. of an ethereal CH₃N₃ soln. prepd. from 8.3 cc. of nitroso-methylurethan. In order to secure complete methylation the operation must be repeated several times, the solvent being distd. off each time. In all, 785 cc. of a 1.5% CH₃N₃ soln. was used. The tetramethylpyrrolone so prepd. m. 219-21° (from MeOH) and formed white needles with a yellowish tint which were highly hygroscopic. By hydrolysis by boiling with dil. HCl, 80% yielded 0.7, 0.7, 0.4 tetramethylolacetone in 105 fl°

M. G. Mouri

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<p>The chemical constitution of hormones stimulating the growth of plants. Zofia Jerzmanowska. <i>Prague Chem. J.</i> 1961 7(11038).—Review. K. Jurek</p>																																																			

thermal decomposition of a few glucosides. Z. Jerzmanowska and Stefania Kłosówna. *Koczniki Chem.* 18, 234-45 (in German, 243-4) (1938).—Ac derivs. of glucosides decompose on heating under reduced pressure, with rupture of the glucosidic bond. The resulting compounds are acetylated aglucons having a free HO group and an unsatd. anhydro sugar. In this way, triacetylhydroxy-2-rhamnal, m. 74°, was obtained from quercitrin. In the case of phlorizin, the products of decompn. undergo further transformations, resulting in derivs. of phloretin and *D*-pentaacetylglucose. M. Wojciechowski

M. Wojciechowski

AS 0-564 METALLURGICAL LITERATURE CLASSIFICATION

ca 10

Condensation of unsaturated esters with urea. *Z. Jergumawaka and I. Gamota. Rozwiti Chem. 18: 2458 (in German, 248 0) (1980).* - By condensation of tetra-*R* ester of ethylenetetracarboxylic acid (I) with urea, thiohydric acid (2,2'-dithio-*A,A'*-bithiuric acid) is obtained. The mechanism of the reaction is explained in the following way. At the first stage I is reduced by Na ethoxide and then the proper condensation takes place with the formation of barbituric ring. M W

ASB.55A METALLURGICAL LITERATURE CLASSIFICATION

DATE	CLASS	NUMBER	ISSUE	YEAR	MONTH	DAY	TIME	DATE	CLASS	NUMBER	ISSUE	YEAR	MONTH	DAY	TIME
10	10	10	10	10	10	10	10	10	10	10	10	10	10	10	10

17

Chem 17

Ergot alkaloids. Zofia Jerzmanowska (Univ. Lodz, Poland). *Wiedomości Chem.* 5, 81-101(1949).—A review with 11 references. Adam Spaszyński

1957

Country : Poland
Category :

G-3

Abs. Jour :

45987

Author : Jerzmanowska, Z. and Skulski, J.
Institut. : Not given
Title : Dimerization and Polymerization Reactions of
Euparin

Orig Pub. : Roczniki Chem, 32, No 3, 471-483 (1951)

Abstract : The substance (I) formed by the dimerization of euparin (2-isopropenylcoumarone) (II) in dil alcoholic solutions of mineral acids (see preceding communication, RZhKhim, No 23, 1955, 55256), absorbs 3 mols H_2 on hydrogenation over Pt (from PtO_2) in glacial CH_3COOH and adds 3 mols perbenzoic acid in chloroform solution. On refluxing for 19 hrs in C_6H_6 with maleic anhydride, I forms the adduct $C_{30}H_{26}O_9$, mp 120° , later 230° [sic] (decomp; from benzene), and on

Card: 1/3

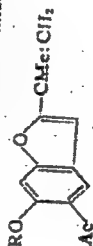
JERZMANOWSKA, ZOFIA

Analiza jakosciowa zwiazkow organicznych; podrecznik laboratoryjny. Warszawa, Panstwowy Zaklad Wydawn. Lekarskich, 1951. 291 p. / Qualitative analysis of organic compounds; a laboratory manual. Index, tables /

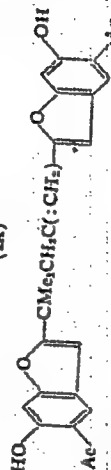
SO: Monthly List of East European Accessions Vol 2 No 9 Library of Congress Sept 53 ~~XXXX~~ Uncl

JERZMANOWSKA, Z.

1/2
Eupatin, the constituent of *Eupatorium cannabinum*, and some derivatives. Z. Jerzmanowska (Med. Akad. Łódź), *Polish Acad. Sci., Pol. Kom. Farm., Warszawa*, Pharm. 3, 165-82 (1951) (English summary). — Euparin (I) (LA, R = H) similar to that isolated by Robertson from American *Eupatorium purpureum*, was isolated from European *E. cannabinum*; however, Robertson's 1 in. 118° and its oxime m. 147° while 1 from *E. cannabinum* m. 121-3° and its oxime m. 150°. Dry. powd. roots (300 g.) were extd. twice with 2.5 l. 80% EtOH for 1.5-2 hrs., the ext. vapd. in vacuo on a water bath to 300 cc., extd. 3 times with Et₂O, the ext. treated with active C, the ether removed, the residue cooled and washed with cold alc. gave 2.4-4.6 g. crude product, which, crystd. from alc., yielded 2.3-2.6 g. 1, thick yellow needles, m. 121-2°, insol. in H₂O and alkali, giving a green-blue coloration with FeCl₃ in alc. soln. To 2.0 g. 1, in freshly distd. quinoline at 30-40° was added 4.5 g. α-bromotetraacetylglucose, then 1.33 g. active Ag₂O, the mixt. mechanically



(IA)



(III)

agitated 10 hrs., 30 cc. concd. AcOH added, the inorg. residue filtered off, the filtrate washed several times with concd. AcOH until almost no turbidity was obtained, and the filtrate added to 500 cc. H₂O giving after 1 hr. 3.5 g. of

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crude cryst. product, which, after several crystallizations from alc. (with C) yielded 1.36 g. (37%) *euparin tetraacetylglucoside* (II) (IA, R = tetraacetyl- α -glucosyl), colorless needles, m. 103-4°, insol. in H₂O, Et₂O, sol. in C₆H₆, CHCl₃, and dioxane. Hydrolysis of 0.07 g. I with 4 cc. of 2.0% alc. HCl gave 0.03 g. of a residue, m. 107-9°, or 0.1 g. I boiled 6 hrs. with 4% alc. H₂SO₄ and let stand 12 hrs. yielded 0.035 g. cryst. yellow residue, m. 107-9° (from alc.); both products, crystallized twice from alc., yielded *euparin dimer* (III), m. 173-4°. III, m. 173° (from alc.) (0.25 g.), was also obtained directly from 1 g. I by boiling 9 hrs. with 120 cc. of 4% alc. H₂SO₄. III is sol. in H₂O, C₆H₆, Et₂O, CCl₄, glacial AcOH, CHCl₃, and insol. in alkali; it gives an emerald-green coloration with alc. FeCl₃. III (0.05 g.) in 15 cc. alc., heated to boiling, and treated hot with a soln. of 2,4-(O₂N)₂C₆H₃NHNH₂ gave 0.055 g. I 2,4-dinitrophenylhydrazone, m. 253-4°, dark red cryst. residue. III (0.25 g.) in 12 cc. Me₂CO treated hot with 0.8 g. K₂CO₃ and 5 cc. MeI and heated 7 hrs. gave 0.3 g. crude product, which sepd. from glacial AcOH in crystals m. 127-8°, giving a green-blue coloration with FeCl₃, and sol. in CHCl₃; apparently, methylation of I was not complete. III (0.1 g.) treated with 1.5 cc. alc., then with 2 cc. 20% KOH and with freshly distil. Me₂SO, gave an exothermic reaction; the product was then heated 1 hr. on a water bath, treated with 5 ml. H₂O, and the residue recrystd. from 3 cc. alc. to yield 0.05 g. cryst. Me deriv. of I, m. 157°, giving a neg. reaction with FeCl₃.

Gene A. Wozniak

JERZMANOWSKA Z.

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547.588.2/4 : 547.483.2 : 547.495.2 : 542.05

Jerzmanowska Z., Jaworska-Królikowska M. Studies on α , β -Unsaturated Carboxylic Acids. Some Reactions of Triethyl- β -Phenylethylene- α , α , β -Tricarboxylate.

C.11

"Studia nad kwasami α , β -nienasyconymi. Niektóre reakcje β -fenyloetyleno- α , α , β -trójkarboksylanu etylowego". Roczniki Chemii (PAN). No. 2, 1934, pp. 417-422.

The condensation was investigated of triethyl phenylethylene- and -phenylethane tricarboxylate with urea in heated sodium ethylate solution. In both cases, there was formation of the same compound (in far better yield in the case of the saturated ester) to which was attributed the formation of 5(α -carbamidobenzyl)-barbituric acid.

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JERZMANOWSKA, Z.

POLON

~~Triterpenes. Zofia Jerzmanowska (Akad. Med. Edg.
Poland). Wiadomosci Chem. 8: 98 (1984). — Classifi-
cation of triterpenes and stereochemistry of pentacyclic
triterpenes are reviewed in some detail. 42 references.~~
Adam Skrzyński

NB
2/21

JERZMANOWSKA, Zofia

Studies on *Eupatorium cannabinum*. II. Isolation of triterpene taraxasterol (α -lactuceryl). Jadwiga Grzybowska, Zofia Jerzmanowska, and Henryk Witkowski. *Roczniki Chem.* 28: 197-212 (1954) (English summary); cf. *C.A.* 48, 5948k. α -Lactuceryl is obtained bound with palmitic acid from petr. ether extn. of the dried flowers, followed by alk. hydrolysis of the residue. A method of obtaining the alc. in 0.8% yield on the wt. of the dried flowers is described. III. Chemical analysis of roots and stems with leaves. Jadwiga Grzybowska and Zofia Jerzmanowska. *Ibid.* 213-31. A chem. analysis of the roots, stems, and leaves is given. The petr. ether extn. of the roots gave, besides euparin, m. 162-3°, α -eupaterol acetate (0.01%), which on hydrolysis gives β -eupaterol, m. 130-7°. The residue from the alc. ext. on alk. hydrolysis gave a probable phytosterol, β -eupaterol, m. 159-62°, and the acetate of $C_{27}H_{48}O$ or $C_{27}H_{46}O$, m. 129-31° (0.002%). Palmitic acid was isolated and the presence of oleic and linolenic acids is established. The petr. ether ext. of the stems and leaves gives α -lactuceryl, m. 218-21° (0.00%). Flavone glucoside which hydrolyzed to quercetin, glycerol, and rhamnose was found. Rutin, m. 187-88°, was found in the stems only (0.4%). Chester Plunk

(2)

SIREMACHKA, Z.; GRZYBOWSKA, J.

"Studies of Eupatorium Cannabinum. Pt. 3. Chemical Analysis of "Cots and Stems with Leaves", P. 213, (ROZNIKI CHEMII, Vol. 28, No. 2, 1954, Warsaw, Poland)

SO: Monthly List of East European Accessions (EFAL), LG, Vol. 4, No. 3, March 1955, Uncl.

JERZMANOWSKA, ZOFIA

α,β-Unsaturated acids. V. The synthesis of *β*-phenyl-ethyl-*α,β*-tricarboxylic acid and its derivatives. Zofia Jerzmanowska and Maria Jaworska-Królikowska (Odra, Wrocław, Poland). *Roczniki Chem.* 28, 397-416 (1954) (English summary); cf. *C.A.* 38, 3335. --BzCOEt (4.5 g.), 4 g. $\text{CH}_2(\text{CO}_2\text{Et})_2$ and 13.5 ml. of a catalyst (7.5 g. anhyd. ZnCl_2 and 15 g. Ac_2O) heated on a steam bath, extd. with 50 ml. Et_2O , washed with water, dried over Na_2SO_4 yields 3.5-7.5 g. $\text{Ph}(\text{EtO}_2\text{C})\text{C}(\text{CO}_2\text{Et})_2$ (I), bp 208-9°, n_D^{20} 1.5116, d_4^{20} 1.1373. Shaking 5 g. I, 15 g. Ba-

$(\text{OH})_2 \cdot 8\text{H}_2\text{O}$ in 150 ml. H_2O for 7 hrs. yields 6.5 g. Ba-salt, which after treating with excess 10% HCl , evap. *in vacuo*, extg. with Et_2O and crystg. from 10% HCl yields 1.2 g. $\text{Ph}(\text{HO}_2\text{C})\text{C}(\text{CO}_2\text{H})_2$ (II), m. 105-7°. Boiling 5.0 g. I with 10% alc. KOH, treating with HCl , extg. with Et_2O and distg. yields 2.5 g. $\text{PhC}(\text{CO}_2\text{Et})\text{CO}_2\text{O.CO}$ (III), m.

84-5°. Heating I for 7 hrs. with KOH yields mono K-salt of II which on treatment with HCl gas in C_6H_6 and crystn. from C_6H_6 yields free acid (IV) of III, m. 131-2°. IV with PCl_5 yields a mixt. of acid chlorides (V) of IV and $\text{PhC}(\text{CO}_2\text{O.CO}_2\text{O.CO})$ (VI). Treatment of this mixt. with PhNH_2

in the cold yields $\text{Ph}(\text{OC})\text{C}(\text{CO}_2\text{NHPh})\text{CO}_2\text{NHPh}$ (VIIa) ($\text{R} = \text{PhNHCO}$, $\text{R}' = \text{Ph}$, VII) m. 121-3° and VIIb ($\text{R} = \text{PhNH}$, $\text{R}' = \text{Ph}$) (VIII), m. 207-8°. Treatment of I with SOCl_2 gives V, which yields VII with PhNH_2 . V and EtOH yields III. IV (0.22 g.) and 0.125 g. PhNH_2 in 2 ml. Et_2O in the cold yield 0.4 g. $\text{Ph}(\text{HO}_2\text{C})\text{C}(\text{CONHPh})\text{CO}_2\text{NHPh}$ (IX) (decomp.). Heating IV (0.4 g.) and 0.83 g. PhNH_2 for 2 hrs. gives the acid analog of VIII ($\text{R} = \text{PhNHCO}$, $\text{R}' = \text{H}$) (X) from which $\text{Ph}(\text{HO}_2\text{C})\text{C}(\text{CONHPh})\text{CO}_2\text{NHPh}$ (X) m. 107-10°, is obtained by treating IX with concd. HCl . On drying *in vacuo* at 80° or by melting II, X gives the amide of IV, m. 167-8°. VI. Some reactions of triethyl *β*-phenyl-ethylene-*α,α,β*-tricarboxylate, III, 417-21 (English summary).—Condensation of 10 g. I with 7.5 g. H_2NCONH_2 in 60 ml. EtOH with 2.99 g. Na yields 0.18 g. $\text{Ph}(\text{CH}_2\text{NCO})\text{CHCH}(\text{CO}_2\text{NH})\text{CO}_2\text{NH.CO}$ (XII). A similar treatment

10 g. $\text{Ph}(\text{EtO}_2\text{C})\text{CHCH}(\text{CO}_2\text{Et})_2$ gives 2 g. XII.

E. Dortschke.

JERZMANOWSKA, Z.

Phytochemical problems. Acta Poloniae pharm. 11 Suppl.:65-66 1955.

1. Zakład Chemii Organicznej A. M., Lodz.
(PLANTS,
medicinal)

JERZANOWSKA, Z.
JERZMANOWSKA, Z.

B. Bobranski's Analiza llosciowa zwiazkow organicznych (A Quantitative Analysis of Organic Compounds); a book review.

p. 415 (Wiadomosci Chemiczne) Vol. 11, no. 7, July, 1957, Wroclaw, Poland

SO: MONTHLY INDEX OF EAST EUROPEAN ACCESSIONS (EEAI) LC, VOL. 7, NO. 1, JAN. 1958

JERZMANOWSKA, Z.
POLAND / Organic Chemistry. Natural Substances and
their Synthetic Analogues. G

Abs Jour: Ref Zhur-Khimiya, No 20, 1958, 67682.

Author : Jerzmanowska Z., Crzybowska J.

Inst : Not given.

Title : Flavonoids from Flowers of the Helichrysum Arenarium.

Orig Pub: Acta polon. phamac., 1958, 15, No 1, 13-14.

Abstract: The information pertaining to the flavodines found in flowers of the Helichrysum arenarium were verified and expanded. "Glycoside A" (I) with a melting point of 152-154° (anhydrous), [α]22D-125° (with 1, alcohol). It may be represented by an empirical formula $C_{21}H_{22}O_{10} \cdot 2H_2O$. It is considered a salipurposide [sic], i.e. a 5- β -D-glycoside of naringenin. An acetate of I

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POLAND / Organic Chemistry. Natural Substances and
their Synthetic Analogues.

G

Abs Jour: Ref Zhur-Khimiya, No 20, 1958, 67682.

Abstract: has 174-177° melting point. Hydrolysis of I causes formation of "substance B" which is naringenin (II). Upon addition of a methyl group to II and I (by using CH_3N_2), followed by hydrolyses, 7,4'-dimethoxy-5-oxyflavanone is produced. "Substance D" probably is apigenin, while "Substance E" probably is a kempherol. Acetylation of the latter results in the formation of 3,7,4'-triace-toxy-5-oxyflavon.

Card 2/2

S/081/63/000/004/021/051
B187/E208

AUTHORS: Jerzmanowska, Zofia, Pijewska, Lucyna

TITLE: On the condensation reaction of phenyl pyruvic acid with ethyl malonate

PERIODICAL: Referativnyy zhurnal. Khimiya, no. 4, 1963, 249 - 250,
abstract 42h153 (Roczn. chem., v. 36, no. 4, 1962, 653 - 663
[Pol.; summaries in Russ. and Eng.] }

TEXT: If unsaturated tricarboxylic acid is to be obtained by reaction of $C_6H_5CH_2COCO_2H$ (I) with $CH_2(COOC_2H_5)_2$ (II) in the presence of a catalyst (CAT) consisting of 1 part $ZnCl_2$ and 2 parts $(CH_3CO)_2O$, the anhydride of 4-acetoxy-naphthalene-2,3-dicarboxylic acid (III; IV acid) along with some $C_6H_5CH=C(OCOCH_3)CO_2H$ (V) are formed but the expected $C_6H_5COCH_2C(CO_2H)=C(COOC_2H_5)_2$ (VI) is not obtained. If there are traces of water in the reaction mass the anhydride of the 4-hydroxy-naphthalene-2,3-dicarboxylic acid (VII; VIII acid) and the ethyl ester of the 7-hydroxy-7,8-dehydronaphthalene-1-carboxylic (or-2-carboxylic acid (IX)) are formed. Attempts

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On the condensation reaction of ...

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to synthesize VI in accordance with Knöwenagel's reaction in alkaline medium (pyridine, alcohol, KOH) were also unsuccessful. VII is converted to III by acetylation. Substance III is rather unstable. Its hydrolysis under very mild conditions gives VII again. Hydrolysis of III under more rigorous conditions, as well as boiling of VII with water, give 4-hydroxy-naphthoic-2 acid (X). Under the action of a catalytic amount of CH_3ONa in CH_3OH III is converted to the methyl ester of X. Reaction of III with $\text{N}_2\text{H}_4 \cdot \text{H}_2\text{O}$ (XI) or $\text{C}_6\text{H}_5\text{NHNH}_2$ (XII) in glacial CH_3COOH gives corresponding derivatives of naphthalazine-2,3 (XIII) and of naphthalazine-2,3-dione-1,4 (XIV). 10.3 g NaHCO_3 in 30 ml water are added to 20 g I, m.p. $150 - 151^\circ\text{C}$, in 70 ml alcohol; the precipitated Na-salt of I (Ia) is dried; 20 g of the latter, 27 g II and 42 ml CAT are boiled for 3 hrs, 100 ml ether are added, washed with water until no more Cl^- ions are present; the precipitate is washed with ether, 20.8 % III, $\text{C}_{14}\text{H}_8\text{O}_3$, m.p. $205 - 206^\circ\text{C}$ (from benzene) is obtained. 4.5 g V, $\text{C}_{11}\text{H}_{10}\text{O}_4$, m.p. $171 - 172^\circ\text{C}$ (from benzene-acetone) is separated from the ethereal mother lye. 20 g undried Ia is boiled with 27 g II in 50 ml CAT for 2.5 hrs, 100 ml ether

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On the condensation reaction of ...

are added, washed with water; 4.35 % VII, $C_{12}H_{16}O_4$, m.p. $241 - 243^{\circ}C$ (from dioxane) are separated from the organic layer. After separation of VII the filtrate is evaporated in vacuo to the half of the initial volume; (temperature of the bath up to $135^{\circ}C$); 1.1 g IX, $C_{15}H_{12}O_3$, m.p. $172-174^{\circ}C$ (from CCl_4 -acetone) results. 0.05 g VII is boiled in 1 ml $(CH_3CO)_2O$ for 5 min, 0.04 g III separates after cooling; 0.5 g III is boiled in 10 ml diluted HCl (1:1) for 1 hr, after cooling 97.3 % VIII, $C_{11}H_{16}O_3$, m.p. $224 - 225^{\circ}C$ (from water) is obtained. 15 ml VII are boiled in 1 ml water for about 15 min (until complete dissolution occurs), and after cooling VIII is obtained. 1 ml 0.1 N CH_3ONa is added to 0.2 g III in 2 ml absolute CH_3OH and 18 ml anhydrous $CHCl_3$, after 2 hrs at $\sim 20^{\circ}C$ it is acidified with 20 % CH_3COOH , the solvent evaporated at $\sim 20^{\circ}C$, and 0.15 g methyl ester of VIII, $C_{12}H_{10}O_3$, m.p. $138 - 160^{\circ}C$ (from water) is obtained. 0.3 g III is dissolved in 10 ml hot alcohol, cooled, the solvent evaporated, the precipitate washed with C_6H_6 . 68.6 % monoethyl ester of IV, $C_{16}H_{14}O_6$

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On the condensation reaction of ...

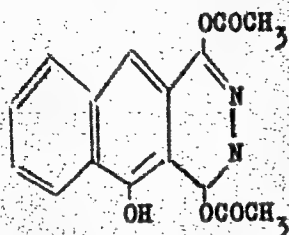
m.p. $143 - 145^{\circ}\text{C}$ (decompos.; from CCl_4), is obtained. 0.5 g III and 0.08 g urea are heated at 200°C for 40 min, 4 ml water are added, 85.7 % imide of VIII, $\text{C}_{12}\text{H}_7\text{NO}_3$, m.p. $295 - 296^{\circ}\text{C}$, is separated (from dioxane). 0.3 g III is mixed with 1 ml 22 % NH_4OH ; after 2 min 31.3 % diamide of IV, $\text{C}_{14}\text{H}_{12}\text{N}_2\text{O}_4$, m.p. $191 - 192^{\circ}\text{C}$, is separated at $\sim 20^{\circ}\text{C}$ (decompos.; from water). 0.5 g III are mixed with 0.5 ml $\text{C}_6\text{H}_5\text{NH}_2$, dissolved in hot acetone after ~ 15 min, by adding C_6H_6 , 63.5 % of the phenylimide of VIII, $\text{C}_{18}\text{H}_{16}\text{NO}_3$, m.p. $167 - 168^{\circ}\text{C}$ (from acetone-benzene) are obtained. 0.5 g III are mixed with 0.15 g $\text{NH}_2\text{OH}\cdot\text{HCl}$ and 0.2 g NaHCO_3 with 2 ml water and 2 ml alcohol, acidified with CH_3COOH after 0.5 hrs at $\sim 20^{\circ}\text{C}$, 38.8 % monooxime of III, $\text{C}_{14}\text{H}_9\text{NO}_5$, m.p. $253 - 254^{\circ}\text{C}$ (decompos.) is obtained. 1 g III is dissolved in 20 ml hot glacial CH_3COOH , 6 drops of about 85 % XI are added, this is boiled for 2 hrs, and after cooling 0.86 g XIII, $\text{C}_{16}\text{H}_{12}\text{N}_2\text{O}_5$, m.p. $258 - 260^{\circ}\text{C}$ (from CH_3COOH -dioxane-water) is separated. In an analogous

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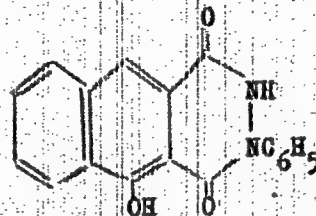
On the condensation reaction of ...

way 50 % XIV, $C_{18}H_{12}N_2O_3$, m.p. $254 - 255^{\circ}C$ (from alcohol) is obtained from 1 g III in 25 ml glacial CH_3COOH and 9 drops XII (boiling for 4 hrs).



XIII

[Abstracter's note: Complete translation.]



XIV

Card 5/5

BARTOSZEWSKI, Jan, dr.; JERZMANOWSKA, Zofia

Condensation of symmetrical diarylthiourea with chloroacetone;
synthesis of new derivatives of thiazoline-4. Pt.1. Roczniki chemii
37 no.1:11-29 '63.

1. Department of Organic Chemistry, Faculty of Pharmacy, Medical
Academy, Lodz.

JURGMANOWSKA, Zofia, prof. dr

Instead of an answer. Problemy 19 (t.c.20) no. 3:130-131 '62.

1. Prorector, School of Medicine, Lodz, Head, Department of
Organic Chemistry, Division of Pharmacy, School of Medicine,
Lodz.

JERZMANOWSKI, Antoni; UBYSZ-JERZMANOWSKA, Krystyna

Variability of *Corynebacterium diphtheriae* following exposure to antibiotics. I. Changes following exposure to penicillin and aureomycin. Med. dosw. mikrob. 10 no.2:193-204 1958.

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(*CORYNEBACTERIUM DIPHTHERIAE*, effect of drugs on,
chlortetracycline & penicillin, variability (Pol))

(*CHLORTETRACYCLINE*, effects,
Corynebacterium diphtheriae variability (Pol))

(*PENICILLIN*, effects,
same)

JERZMANOWSKI, Antoni; UBYSZ-JERZMANOWSKA, Krystyna

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(*CORYNEBACTERIUM DIPHTHERIAE*, pharmacol.)

(*STREPTOMYCIN*, pharmacol.)

(*CHLORAMPHENICOL*, pharmacol.)

(*OXYTETRACYCLINE*, pharmacol.)

ONISK, Zbigniew; JERZMANOWSKI, Antoni

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(DERMATOLOGY) (STAPHYLOCOCCAL INFECTIONS)

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p. 177.

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May, 1959, Unclass.

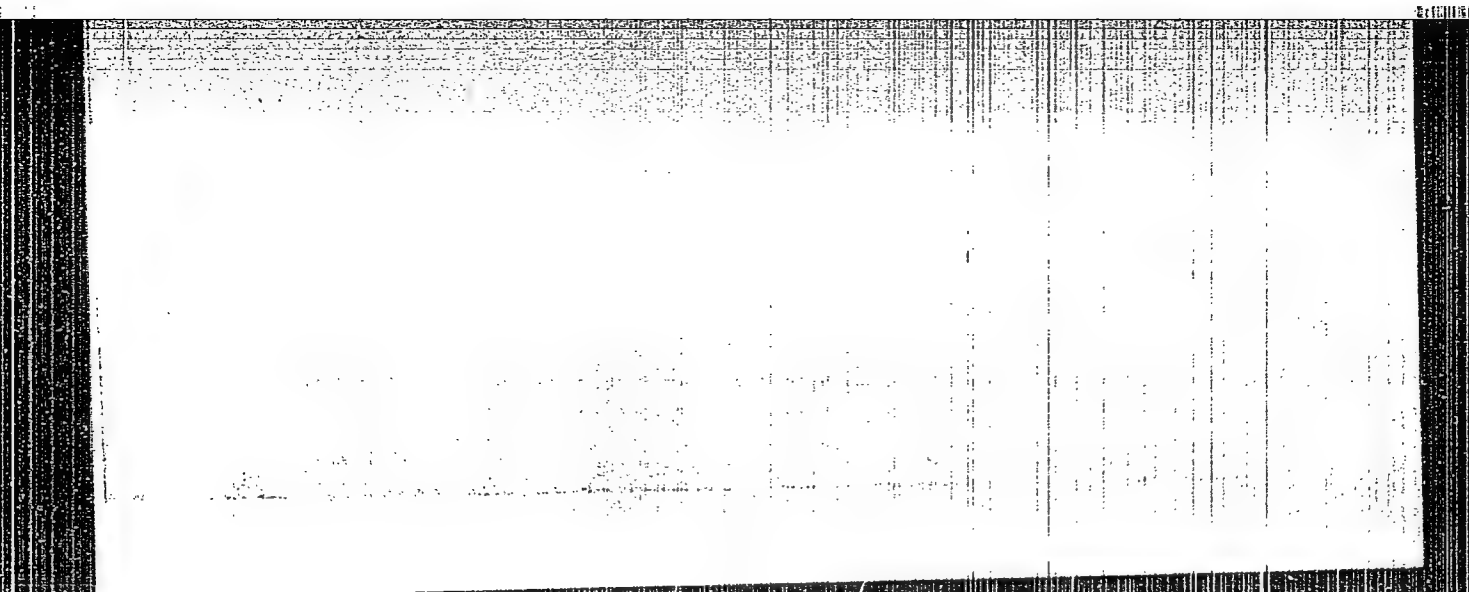
JERZMANSKA, Anna; JUCHA, Stanislaw

Significance of the ichthyofauna in schists of Jaslo near Dynow.
Rocz geol Krakow 33 no.1/3:159-180 '63.

1. Katedra Paleozoologii, Uniwersytet, Wroclaw, i Katedra
Geologii, Akademia Gorniczo-Hutnicza, Krakow.

"APPROVED FOR RELEASE: 08/10/2001

CIA-RDP86-00513R000619620004-3



APPROVED FOR RELEASE: 08/10/2001

CIA-RDP86-00513R000619620004-3"

JERZMANSKI, Jerzy

POLAND

JERZMANSKI, Jerzy

Lower Silesian Field Station, Geological Institute
(Dolnoslaska Stacja Terenowa Instytutu Geologicznego)

Warsaw, Kwartalnik geologiczny, No 3, 1963, pp 529-30.

"Main Directions in Iron Ore Deposit Explorations in
the Sudetes".

JERZYANSKI, Jerzy

Problems of prospecting for sedimentary iron ore deposits in
the Caledonian series of the Sudetes. Przegl geol. 11 no.9:
410-413 S'63

JERZYKIEWICZ, A

621.313.616.9 : 621.314.21.018
 ✓ 4123. The possibility of applying synthetic materials
 as high-voltage insulation in Polish-made testing
 transformers. A. JERZYKIEWICZ AND M. KULAK.
 Przeglad elektrotechniczny, Warszawa, 245-9 (1953) in
 Polish.
 The difficulties encountered in the production of
 dry h.v. transformers are described. They may be
 used only up to 30 kV. Synthetic materials are
 discussed and special requirements for voltage trans-
 formers (with one earthed or both terminals un-
 earthed) and current transformers (high mechanical
 strength of insulation on short-circuit) are mentioned.
 Examples of transformers with plastic insulation are
 given.
 N. W. MAKOWSKI

JERZYKIEWICZ A

POLAND/Chemical Technology - Chemical Products and Their
Application - Synthetic Polymers. Plastics.

H.

Abs Jour : Ref Zhur - Khimiya, No 9, 1958, 30319
Author : Hertz, Z. and Jerzykiewicz, A.
Inst : -
Title : Epoxide Resins in the Electrical Industry.
Orig Pub : Przegląd Elektrotechn, 31, No 10-11, 641-643, 1955.
Abstract : See RZhKhim, 1957, 9709.

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2883

876.24.01.013

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"Urządzenia do wymiany sita", Przegląd Papierniczy, No. 3, 1953, pp. 67-71, 14 figs.

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JERZYKIEWICZ, J.

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SO: Monthly List of East European Accessions. (EEAL). LC. Vol. 4, No. 4.
April 1955. Uncl.

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SO: Monthly List of East European Accessions. (NEAL). LC. Vol. 4, No. 4. April 1955. Uncl.

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SO: Monthly List of East European Accessions (EEAL) LC. Vol. 6, No. 12, Dec. 1957.
Uncl.

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A Polish made paper machine put into operation in Turkey. Przegl
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1. Fabryka Maszyn Papierniczych, Cieplice (for Jerzykiewicz).
2. Biuro Projektow Przemyslu Papierniczego, Lodz (for Anglik).

JERZYKIEWICZ, M.

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no.4:253-260 '63.

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KOCHANOWICZ, Teresa; JERZYKOWSKA, Halina

Treatment of recurrent aphthae with atabrine. Przegl.derm.,
Warsz.46 no.5:489-493 8-0 '59.

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prof.dr. T. Chorazak.
(QUINAGRINE ther.)
(STOMATITIS ther.)

JERZYKOWSKA-KULESZYNA, K.

Two cases of exchange blood transfusion in newborn with hemolytic disease. *Pediat. polska* 26 no.2:200-203 Feb 1951. (CIML 21:1)

1. Of the Obstetric-Gynecological Clinic (Director -- Prof. T. Zwolinski, M.D.) and of the Pediatric Clinic (Director -- Prof. K. Jonscher, M.D.) of Poznan Medical Academy.

JERZYKOWSKA-KULESZYNA, K.; ZYWICKA-TWAROWSKA.

Mortality of newborn infants in 1951; data of the Newborn Ward of the Obstetric and Gynecologic Clinic of the A. M. W. Poznan. *Pediat. polska* 27 no. 9:1091-1097 Sept 1952. (CLML 23:3)

1. Of the Newborn Ward (Head —Prof. K. Jonscher, M. D.) of Obstetric-Gynecological Clinic (Director—Prof. I. Roszkowski, M.D.) of Poznan Medical Academy.

JERZYKOWSKA-KULESZYNA, K.

Exchange blood transfusion in fetal erythroblastosis. *Pediat. polska* 27 no. 11:1325-1338 Nov 1952. (GLML 24:1)

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1. Z Kliniki Położniczo-Ginekologicznej Akademii Medycznej w Poznaniu; kierownik: prof. dr I. Reszkowski; kierownik: naukowy Oddziału Noworodków; prof. dr K. Jenzsher i z Zakładu Medycyny Sądowej Akad. Medycznej w Poznaniu; p.o. kierownika: dr. E. Chroscielewski. Poznań ul. Świecickiego 6. Zakład Medycyny Sądowej A.M.

(LUNGS, diseases,
hyaline membrane)
(INFANT, NEWBORN, diseases,
pulm.hyaline membrane)

JERZYKOWSKA-KULESZYNA, Kazimiera.

Observation on the therapy of hemolytic disease in newborn with exchange transfusion of blood. *Pediat.polska* 30 no.2:139-150 Feb.'55.

Z Gdzialu Noworodkow Kliniki Poloznictwa i Chorob Kobiacych A. M. w Poznaniu. Kierownik Kliniki: prof. dr med. I. Roszkowski, Konsultant naukowy oddz. noworodkow: prof. dr med. K. Jonscher Adres Poznan, Matejki 6.

(BLOOD TRANSFUSION

exchange, in ther. of hemolytic dis. in newborn inf. indic.)

(ERYTHROBLASTOSIS, FETAL, therapy

blood transfusion, exchange, indic.)

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A.M. Poznaniu. Kierownik: prof. dr. med. I. Roszkowski. Kon-
sul'tant oddzialu noworodkow: prof. dr med. K. Jonsher. Poznan,
Polna 33.

(ERYTHROBLASTOSIS, FETAL, therapy,
progn.)

POLAND/Pharmacology. Toxicology. Tranquilizers.

V

Abs Jour: Ref. Zhur. - Eiol., No 22, 1958, 102781

Author : Jerzykowska-Kuleszyna, Kazimiera

Inst : ~~Z Oddzialu Noworodkow i Klinika Położnictwa i Chorob Kobietych A.M. w~~
Boznaniu. Kierownik: doc dr Med. W. Michalkiewicz. Kierownik Oddzialu:
doc dr med. K. Jerzykowska-Kuleszyna.

Title : The Application of Largactyl in Diseases of the
Newborn, Including Premature Babies.

Orig Pub: Ginekol. Polska, 1958, 29, No. 1, 33-41

Abstract: Largactyl (I; 2 mg/kg daily) was introduced intra-
muscularly or internally. Beneficial effect of
I in cases of birth injury of babies born at term
was noted; in premature babies I is less effective.
In a group of newborn which were subjected to sur-
gery, I considerably decreased the lethality. In
hemolytic diseases of the newborn, I did not
improve the results of blood transfusion. Bibl.
28 items. - From the author's resume.

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BERNARDCZYKOWA, Anna; JERZYKOWSKA-KULESZYNA, Kazimiera

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A. Kwaskowski Z Oddziału Noworodków i Wczesniaków I Kliniki Położnictwa
i Chorob Kobietych A. M. w Poznaniu. Kierownik: doc. dr med. W.
Michalkiewicz. Adres: dr Anna Bernardczykowa, Poznań, ul. Dwierczewskiego
31 m. 11.

(RETROLENTAL FIBROPLASIA, case reports
(Pol))

JERZYKOWSKA-KULESZYNA, Kazimiera; KRZYWINSKA, Felicja

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Kierownik: doc. dr med. W. Michalkiewicz. Adres: Poznan, ul. Polna 33, I Klinika Poloznicstwa i Chorob Kobietych A.M.

(ACTH, eff.
on adrenal cortex in newborn inf. in normal & pathol. states (I
(ADRENAL CORTEX, eff. of drugs on,
ACTH, in newborn inf. in normal & pathol. states (Pol))
(INFANT, NEWBORN
eff. of ACTH on adrenal cortex in normal & pathol. states (Pol

JERZYKOWSKA-KULESZYNA, K.; RENZ-SOLAWA, M.; ZYWICKA-TWAROWSKA, I.

Comparative evaluation of clinical and radiological lung examinations in newborn infants. *Pediatr pol* 36 no.1:5-13 '61.

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Kierownik: doc. dr med. W. Michalkiewicz i z Zakładu Radiologii
Lekarskiej A.M. w Poznaniu Kierownik: doc. dr med. B. Gładysz.

(LUNG DISEASES in inf & child) (INFANT NEWBORN dis)

EXCERPTA MEDICA Sec 7 Vol 13/2 'Pediatrics Beb 59

377. LARGACTIL IN THE DISEASES OF NEWBORNS AND PREMATURE INFANTS - Largactyl w chorobach noworodków i wcześniaków - Jerzykowska-Kuleszyna K. Odd. Noworodków i Klin. Położnictwa Chor.

Kobleczyński A.M., Poznań - GINEK. POL. 1958, 29/1 (33-41) Tables 1
In 71 newborns and premature infants, largactil was given i.m. or orally in a dose of 2 mg./kg. per 24 hr., mostly during the first 3 days of life. The author observed an advantageous effect in the cases of labour trauma in the mature infants. A decrease of the mortality rate of the newborns submitted to operation could be observed. No beneficial effect was observed in cases with exchange transfusions. Largactil increased the difficulties during loss of blood by a lowering of the blood pressure.
Dziesięszewska - Warsaw (VII, 10)

JERZYKOWSKA, Kazimiera; BREBOROWICZ, Alfreda; SZCZEPANSKA, Zofia;
STULKOWSKI, Kazimierz

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BILINSKI, Zbigniew; JERZYKOWSKI, Mieczyslaw; MATKOWSKI, Jozef

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(BLOOD,

carbon monoxide, photometry)

(CARBON MONOXIDE, in blood,
determ., photometry)

JERZYKOWSKI, Tadeusz

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Rokitnicy: kierownik: doc. dr S. Jozkiewicz)
(HEMOGLOBIN)

JERZYKOWSKI, Tadeusz

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(VITAMIN D chem)

JERZYKOWSKI, Tadeusz

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(Vitamins) (Chemical reactions)

JERZYKOWSKI, Tadeusz

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JES, S. - Inzenyrske Stavby Vol. 3, no. 1, Jan. 1955

Conference on the limiting states of structural elements. p.37

SO: Monthly List of East European Accessions, (EEAL), LC, Vol. 4, No. 9, Sept. 1955, Uncl.

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p. 358.

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Monthly list of East European Accessions (EEAI) LC. Vol. 9, no. 2, Feb. 1960
Uncl.

JES, S.

JES, S. Beams on elastic bases. p. 428.

Vol. 4, No. 9, Sept. 1956.
INZENYRSKE STAVBY.
TECHNOLOGY
Praha, Czechoslovakia,

So: East European Accession, Vol, 6, No. 3, March 1957

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Je '63.

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JESE, Leopold

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(EYE, wds. & inj.
ther.)

(WOUNDS AND INJURIES
eye, ther.)

(EYE, dis.
ther.)

JESE, Leopold, prof. dr.

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prof. dr. L.Jese)
(CATARACT, ther.
(Ser))

JESENAK, J.

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Praha, Czechoslovakia

SOURCE: East European List (EEAL) Library of
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JESENAK, Jan, inz.

Observations on the static formula in the new Czechoslovak
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Modern method of calcination of pyrite. J. Beňa and V. Jesenák (Fac. Inorg. Technol., Bratysłava, Czech.). *Chem. Průmysl* 10, 343-7 (1953). — The "fluidized operation," calcining pyrite powder suspended in a current of air, is discussed. The output of SO₂ is increased, installation costs are low, the temp. is easily maintained, and the grade of pyrite can be varied. This article is from the Czech literature.

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Continuation of report for the period 1/1/68 to 12/31/68
L. A. Howard

APPROVED FOR RELEASE: 08/10/2001

CIA-RDP86-00513R000619620004-3"

Jesenak V

CZECHOSLOVAKIA / Chemical Technology. Chemical Products. H
Processes and Apparatuses of Chemical Technology.

Abs Jour: Ref Zhur-Khimiya, 1958, No 20, 67695.

Author : Kossaczky E., Bena J., Jesenak V., and Ilavsky J.
also Singer D.

Inst : Not given.

Title : Discussion of Singer's Article "Theoretical Bases
of Processes Involving Pseudoliquification" and
Answers to the Discussions by Beranka and Klumper.

Orig Pub: Chem. prumysl, 1956, 6, No 10, 430-433.

Abstract: Ref to Ref. Zhur-Khimiya, 1958, 25349. No abstract.

Card 1/1

L 37751-66 EWP(j)/T DS/RM

ACC NR: AT6028246

SOURCE CODE: HU/2502/65/046/001/0035/0044

AUTHOR: Braun, Tibor (Doctor; Budapest); Hradil, M.—Khradil, M. (Bratislava); ⁴⁴
Jesenak, V.—Yesenak, V. (Bratislava); Tolgyessy, J.—Tel'deshi, Y. (Doctor; Bratislava) ²⁺¹

ORG: [Braun] Institute of Inorganic and Analytical Chemistry, L. Eotvos University, Budapest; [Hradil; Jesenak; Tolgyessy] Department of Radiochemistry and Radiation Chemistry, Faculty of Chemistry, Slovak Technical University, Bratislava, Czechoslovakia ^{1/4}

TITLE: Radiocoulometric titrations

SOURCE: Academia scientiarum hungaricae. Acta chimica, v. 46, no. 1, 1965, 35-44

TOPIC TAGS: titrimetry, radiation chemistry, radioisotope

ABSTRACT: Two methods, one intermittent and the other continuous, have been developed for radiometric determination of the end point of coulometric titrations based on formation of precipitate and complexation. In the radiocoulometric titrations based on precipitate formation, iodide ions labeled with I^{131} were titrated with silver ions generated by coulometry, using silver electrodes. In the complexometric radiocoulometric titrations with the aid of a solid indicator, the cyanide ions generated by the electrolysis of the complex $[Ag(CN)_2]^-$ were reacted with the Ni^{++} ions to be determined, using AgI solution containing labeled Ag. The experimental apparatus is described. Orig. art. has: 5 figures and 1 table. [Orig. art. in Eng.] [JPRS: 33,906]

SUB CODE: 07 / SUBM DATE: 30Jan65 / ORIG REF: 010 / OTH REF: 002

Card 1/20

0425 1651

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inz., CSc.; BRAUN, Tibor, dr.

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M-4

Abs Jour : Ref Zhur - Biol., No 3, 1958, 10925

Author : Jesic, D.

Inst : Institute of Beet Husbandry (Crvenka)

Title : Application of Microelements in the Cultivation of Sugar
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Orig Pub : Poljopr. Vojvod., 1956, 4, No 11, 13-22

Abstract : Investigations into problems of the application of B and
Mg were conducted by the Institute of Beet Husbandry in
the city of Crvenka (Vojvodina, Yugoslavia). When beet
was fertilized with the minimal quantity of B (in the
form of borax) combined with NPK, the yield increased on
the average by 6%. The sugar content increased by 1.5%.

Card 1/2

YUGOSLAVIA/Cultivated Plants - Technical, Oil, and Sugar
Plants.

M-4

APPROVED FOR RELEASE: 08/10/2001

CIA-RDP86-00513R000619620004-3"

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